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# **Regulation of AURORA B function by mitotic checkpoint protein MAD2**

Jayasha Shandilya<sup>1</sup>, Kathryn F Medler<sup>1</sup> and Stefan GE Roberts<sup>1,2\*</sup>

<sup>1</sup>Department of Biological Sciences, University at Buffalo, Buffalo, NY 14260

<sup>2</sup>School of Cellular and Molecular Medicine, University of Bristol, Bristol BS8 1TD, UK

\*Correspondence

Email: sr237@buffalo.edu

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## Abstract

Cell cycle checkpoint signaling stringently regulates chromosome segregation during cell division. MAD2 is one of the key components of the spindle and mitotic checkpoint complex that regulates the fidelity of cell division along with MAD1, CDC20, BUBR1, BUB3 and MAD3. MAD2 ablation leads to erroneous attachment of kinetochore-spindle fibres and defective chromosome separation. A potential role for MAD2 in the regulation of events beyond the spindle and mitotic checkpoints is not clear. Together with active spindle assembly checkpoint signaling, AURORA B kinase activity is essential for chromosome condensation as cells enter mitosis. AURORA B phosphorylates histone H3 at serine 10 and serine 28 to facilitate the formation of condensed metaphase chromosomes. In the absence of functional AURORA B cells escape mitosis despite the presence of misaligned chromosomes. In this study we report that silencing of MAD2 results in a drastic reduction of metaphase-specific histone H3 phosphorylation at serine 10 and serine 28. We demonstrate that this is due to mislocalization of AURORA B in the absence of MAD2. Conversely, overexpression of MAD2 concentrated the localization of AURORA B at the metaphase plate and caused hyper-phosphorylation of histone H3. We find that MAD1 plays a minor role in influencing the MAD2-dependent regulation of AURORA B suggesting that the effects of MAD2 on AURORA B are independent of the spindle checkpoint complex. Our findings reveal that, in addition to its role in checkpoint signaling, MAD2 ensures chromosome stability through the regulation of AURORA B.

## Introduction

MAD2 is a critical member of the spindle assembly checkpoint (SAC) and mitotic checkpoint (MCC) complexes. At the onset of mitosis MAD2 and its interacting partner MAD1 bind to unattached kinetochores and activate SAC signaling to ensure proper spindle microtubule and kinetochore attachments and the fidelity of chromosome segregation<sup>1</sup>. MAD2 together with other checkpoint proteins CDC20, BUBR1, BUB3 and MAD3 form the MCC which blocks the function of Anaphase promoting complex/Cyclosome (APC/C). This pauses the cells in metaphase until all the chromosomes are properly attached and bi-oriented<sup>2,3</sup>.

In addition to SAC- and MCC-based regulation of the cell cycle, several kinases and phosphatases also play important roles in guiding stringent chromosome segregation. AURORA B kinase is a member of the chromosome passenger complex (CPC) which associates with the inner centromeres and regulates chromosome separation during mitosis<sup>4</sup>. AURORA B targets histone H3 S10 and S28 and their phosphorylation marks the condensed metaphase chromosomes<sup>5</sup>. Recent studies have suggested a possible link between MAD2 and AURORA B function during mitosis<sup>6</sup>. However it is not clear how MAD2 directly regulates AURORA B function to promote proper chromosome separation and genomic stability. We show here that MAD2 is a critical mediator of chromosome segregation by regulating the mitotic localization and function of AURORA B. RNAi-mediated silencing of MAD2 results in aberrant localization of AURORA B at the metaphase plate and consequent loss of histone H3 phosphorylation. Our results also show that MAD2 (not MAD1) is the critical SAC component which modulates mitotic functions of AURORA B and is instrumental in preventing chromosome-segregation defects.

## Results

### MAD2 regulates AURORA B-mediated mitotic phosphorylation of histone H3

Our earlier work suggested that MAD2 might play a role in events leading to chromosome condensation during mitosis <sup>7</sup>. To study the effect of MAD2 on epigenetic changes linked to condensed metaphase chromosomes <sup>8, 9</sup> we silenced MAD2 in HeLa cells and found that the phosphorylation status of histone H3 S10 and S28 residues was drastically reduced (Fig. 1A). Immunofluorescence analysis of metaphase chromosomes in HeLa cells revealed mitotic specific reduction of histone H3 phosphorylation after MAD2 silencing (Fig. 1B). RNAi-mediated silencing of AURORA B also resulted in a significant reduction of H3 S10 and S28 phosphorylation levels (Fig. 1A, 1C). We next performed siRNA-rescue experiments to confirm the specificity of the observed effect on H3 phosphorylation. Our results showed that co-transfection of a MAD2 expression plasmid <sup>10</sup> (that is RNAi-resistant in our system) can rescue the phosphorylation of histone H3S28 in MAD2-silenced HeLa cells. The recovery in the level of histone H3 phosphorylation was comparable to the control cells (Fig. 1D).

Similar to the observations above, our recent study showed that MAD2 knockdown in WiT49 and M15 cells also resulted in a drastic reduction in H3 S10 phosphorylation <sup>7</sup>. Interestingly, knockdown of either MAD2 or AURORA B in MCF7 cells had negligible effect on mitotic histone H3 S10 phosphorylation but still caused a significant reduction in phosphorylation of histone H3 (S28) (Supplementary Fig. S1A). Taken together the results suggest cell type specific effects of MAD2 depletion on histone H3 phosphorylation.

At the onset of mitosis, histone H3 is known to be phosphorylated by HASPIN kinase at Threonine 3 residue. Phosphorylation of Histone H3T3 is crucial for the recruitment of

AURORA B at the centromeres<sup>11, 12</sup>. However, AURORA B has been shown to regulate the phosphorylation status of histone H3T3 via a feedback loop<sup>13, 14</sup>. We therefore determined if MAD2 knockdown influences H3T3 phosphorylation. Interestingly, we found that MAD2 ablation resulted in a significant reduction of H3T3 phosphorylation in both HeLa and WiT49 cells (Fig. 1E). This result shows that MAD2 most likely affects multiple signaling pathways by directly regulating AURORA B. Hence silencing MAD2 negatively affects AURORA B mediated phosphorylation of the histone H3 tail which is essential for mitotic chromosomal condensation.

#### **MAD2 regulates AURORA B localization during mitosis**

The expression and activity of AURORA B peaks during mitosis<sup>5</sup>. AURORA B along with three other proteins, SURVIVIN, BOREALIN and INCENP form the chromosome passenger complex (CPC)<sup>4, 15</sup>. Our results so far have shown that MAD2 ablation results in the reduced phosphorylation of histone H3. In order to identify the role of MAD2 in the regulation of mitotic localization of AURORA B, we silenced MAD2 in different cell lines and studied AURORA B localization using immunofluorescence analysis. Our results in HeLa, WiT49 (Figs. 2A and 2B) and MCF7 cells (Supplementary Fig. S1B) show that AURORA B occupancy at the metaphase plate is reduced upon MAD2 silencing.

MAD2 ablation is known to promote early anaphase entry due to inactive mitotic checkpoint function, which might account for the effects that we have observed so far. To rule out this possibility we performed experiments in presence of the proteasome inhibitor MG-132 (5  $\mu$ M) for 4 hours<sup>6</sup> which effectively prevents mitotic exit and arrests the cells at metaphase

(Fig. 2C). The cells were transfected with either MAD2 siRNA or AURORA B siRNA (similar to that described in Fig. 1), followed by a 4 hour treatment with either DMSO or MG-132 (5  $\mu$ M). Our results show that delaying anaphase entry did not significantly change the effect of silencing MAD2 or AURORA B on histone H3 (S28) phosphorylation (Fig. 2D). Taken together, our data show that MAD2 has a profound effect on AURORA B recruitment and function during mitosis and provide strong evidence that AURORA B can be regulated by SAC signaling components. Interestingly, we found no significant defects in the mitotic localization of SURVIVIN (another member of CPC) <sup>4, 15</sup> upon MAD2 silencing in HeLa and WiT49 cells (Figs. 2E and 2F). This observation suggests that MAD2 may not influence the functional integrity of the CPC but rather affects specific properties of AURORA B.

### **MAD2 overexpression enhances AURORA B-mediated phosphorylation of histone H3 during mitosis**

Silencing MAD2 leads to mislocalized AURORA B and reduced histone H3 phosphorylation during mitosis which promotes error-prone cell division. We next tested whether MAD2 overexpression conversely affects AURORA B function <sup>10</sup>. Fig. 3A shows that overexpression of MAD2 increased phosphorylation of histone H3 at both S10 and S28 residues. Moreover, HeLa cells transfected with MAD2 overexpression construct exhibited enhanced AURORA B signal at the metaphase plate when compared to the control transfected cells (Fig. 3B). Again, blocking early mitotic exit using MG-132 (5  $\mu$ M) for 4 hours had no significant effect on the elevation of histone H3 (S28) hyper-phosphorylation in MAD2 overexpressing cells when compared to control DMSO-treated cells.

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## 125 **MAD1 and MAD2 complex regulates mitotic functions of AURORA B**

126       Our results so far have shown that perturbation in MAD2 protein level significantly alters  
127 the status of histone H3 phosphorylation and AURORA B occupancy during mitosis. MAD2  
128 plays an important role in the initiation of spindle assembly checkpoint signaling along with  
129 MAD1 at the kinetochores. MAD1 recruits soluble O-MAD2 (open conformation of MAD2) and  
130 stimulates the conversion to C-MAD2 (closed active conformation of MAD2). During pro-  
131 metaphase the MAD1-MAD2 tetramer binds to the unattached kinetochores and amplifies SAC  
132 signaling<sup>1-3</sup>. In order to determine if MAD1-mediated initial recruitment of MAD2 is the critical  
133 step in regulating AURORA B function, we silenced MAD1 in HeLa cells and analyzed the  
134 phosphorylation levels of histone H3 (S10 and S28). Interestingly, loss of MAD1 resulted in only  
135 a minor reduction in the level of histone H3 phosphorylation (S10 or S28; Fig. 4A), when  
136 compared to the effect of MAD2 silencing (Fig. 1A). Furthermore, immunofluorescence analysis  
137 of MAD1-silenced cells did not show a significant difference in either AURORA B occupancy  
138 or histone H3 phosphorylation during metaphase (Fig. 4B). These observations suggest that the  
139 effects of MAD2 reported here are only marginally influenced by MAD1 and thus soluble C-  
140 MAD2 is the likely entity that regulates the events downstream of the initial SAC activation.

141



## Discussion

The fidelity of chromosome segregation during cell division is controlled by the delicate balance of expression and activity of several spindle and mitotic checkpoint proteins<sup>16</sup>. Loss of MAD2 or other mitotic checkpoint components promotes catastrophic events such as early mitotic exit with chromosome mis-segregation and accumulation of aneuploidy. Activation of checkpoint signaling arrests the cells at metaphase until all the attachment errors are rectified and the chromosomes are properly bi-oriented at the metaphase plate. Together with these checkpoint proteins, the expression of checkpoint-regulatory factors such as p31 comet<sup>17, 18</sup> and recently identified, WT1<sup>7</sup> modulate the signaling pathway via MAD2 interaction and regulate the timing of mitotic exit.

In this study we have demonstrated that silencing of MAD2 has effects beyond spindle/mitotic checkpoint function that are critical for the function of AURORA B kinase, one of the key members of CPC. MAD2 is important for the proper localization of AURORA B and also AURORA B-dependent histone H3 phosphorylation during mitosis. How MAD2 connects with AURORA B is not clear. Our data could be explained either by direct regulation of AURORA B catalytic activity, control of AURORA B localization, or a combination of these effects. The results presented here suggest that the regulation of AURORA B by MAD2 is likely to be independent of the MCC, but whether other intermediary factors are involved will require further studies. Our findings add a new dimension to the role of MAD2 in cell division where it regulates AURORA B recruitment and mitotic phosphorylation of several key residues of histone H3 (T3, S10 and S28). However, cell type specific differences may further fine-tune the effect of MAD2 on AURORA B function as evident from our results with MCF7 cells. Interestingly, an earlier study<sup>6</sup> also reported contrasting effects in the non-cancerous cell line RPE1. It is

165 therefore likely that additional cellular factors play an important role in MAD2-mediated  
166 regulation of AURORA B. In summary, our results suggest that MAD2 acts as a junction in  
167 connecting multiple cell cycle checkpoint signaling pathways (SAC, MCC and CPC) to maintain  
168 genomic stability.

169

## **Materials and Methods**

### **Cell culture and Transfection**

HeLa, WiT49 and MCF7 cells were grown in DMEM supplemented with 10% (vol/vol) FBS at 37°C. Transfection of plasmids was performed using Effectene reagent (Qiagen). Cells were harvested 48 hours after transfection and processed for different assays. MAD2, MAD1 and AURORA B siRNAs were obtained from Qiagen. siRNAs were transfected using Hiperfect reagent (Qiagen) for 48 hours. For the MAD2 rescue experiment siRNA targeting 3'UTR of MAD2 (CTGAAAGTAACTCATAATCTA) was used which has no affect on the coding sequence of the MAD2 overexpression plasmid.

### **Western Blotting and Immunofluorescence Analysis**

Western blotting analysis was performed as described before<sup>7</sup> using antibodies: MAD2 (C-19-sc-6329) and SURVIVIN (D-8, sc-17779) were obtained from Santa Cruz Biotechnology. MAD2 (A300-301A) and MAD1 (A300-339A) antibodies were obtained from Bethyl laboratories. H3S10p (ab5176), H3S28p (ab5169), H3 (ab1791), AURORA B (ab3609) and  $\beta$ -TUBULIN (ab6046) antibodies were obtained from Abcam. H3T3p (07-424) antibody was obtained from Millipore. Immunofluorescence analysis was performed as previously described<sup>7</sup>.

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191 The authors declare no competing financial interests.

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## Figure Legends

### **Figure 1:** MAD2 regulates AURORA B-mediated mitotic phosphorylation of histone H3

(A). HeLa cells were transfected with control, MAD2 siRNA or AURORA B siRNA for 48 hours, followed by immunoblotting with anti-MAD2, anti-AURORA B and anti-TUBULIN antibodies for control. The phosphorylation status of histone H3 residues serine 10 and serine 28 were probed with phospho-site specific antibodies. Histone H3 Western blotting was performed as a control. (B). Immunofluorescence analysis of control and MAD2 siRNA transfected HeLa cells was performed with the specific antibodies to visualize phosphorylation status of histone H3 (S10 and S28) during metaphase. DNA was stained with Hoechst. (C). AURORA B siRNA transfected HeLa cells were analyzed as in B. Scale bar is 10 microns. (D). HeLa cells were transfected with either control or MAD2 siRNA for 48 hours. The control and siRNA transfected cells (24 hours after first transfection) were then co-transfected with either vector or MAD2 expression plasmid (that is RNAi-resistant) for another 24 hours. The cells were then immunoblotted with anti-MAD2, anti-phospho-H3S28 and anti-TUBULIN antibodies. (E). HeLa cells and WiT49 cells were transfected with either control or MAD2 siRNA for 48 hours and Western blotting was done with anti-MAD2, anti-phospho-H3T3 and anti-TUBULIN antibodies.

### **Figure 2:** MAD2 regulates AURORA B localization during mitosis

(A). HeLa and (B) WiT49 cells were transfected with control or MAD2 siRNA for 48 hours, followed by immunofluorescence analysis with anti-MAD2 and anti-AURORA B antibodies to visualize the mitotic localization of AURORA B during mitosis. DNA was stained with Hoechst. Scale bar is 10 microns. (C). Mitotic index showing percentage of HeLa cells in mitosis after

treatment with either DMSO (control) or MG-132 (5 $\mu$ M) for 4 hours (D). HeLa cells were transfected with control, MAD2 and AURORA B siRNA for 48 hours followed by treatment with either DMSO or MG-132 (5  $\mu$ M) for 4 hours. Immunoblotting was performed in with anti-MAD2, anti-AURORA B and anti-TUBULIN antibodies. The phosphorylation status of histone H3 at serine 28 was probed using its phospho-site specific antibody. Histone H3 Western blotting was performed as a control. (E). HeLa cells and (F). WiT49 cells were transfected with control or MAD2 siRNA for 48 hours, followed by immunofluorescence analysis with anti-MAD2 and anti-SURVIVIN antibodies. DNA was stained with Hoechst. Scale bar is 10 microns.

**Figure 3:** MAD2 overexpression enhances AURORA B-mediated phosphorylation of histone H3 during mitosis

(A). HeLa cells were transfected with an empty vector or MAD2 overexpression plasmid for 48 hours, followed by immunoblotting with anti-MAD2 and phosphorylation specific antibodies against histone H3 residues serine 10 and serine 28. Western blotting with histone H3 and TUBULIN antibodies were performed as controls. (B). HeLa cells were transfected with vector or MAD2 overexpression plasmid for 48 hours, followed by immunofluorescence analysis with anti-MAD2 and anti-AURORA B antibodies to visualize the mitotic localization of AURORA B during metaphase. DNA was stained with Hoechst. Scale bar is 10 microns. (C). HeLa cells were transfected with vector or MAD2 overexpression plasmid for 48 hours followed by treatment with either DMSO or MG-132 (5  $\mu$ M) for 4 hours. Immunoblotting was performed in with anti-MAD2 and anti-AURORA antibodies. The phosphorylation status of histone H3 at serine 28 was

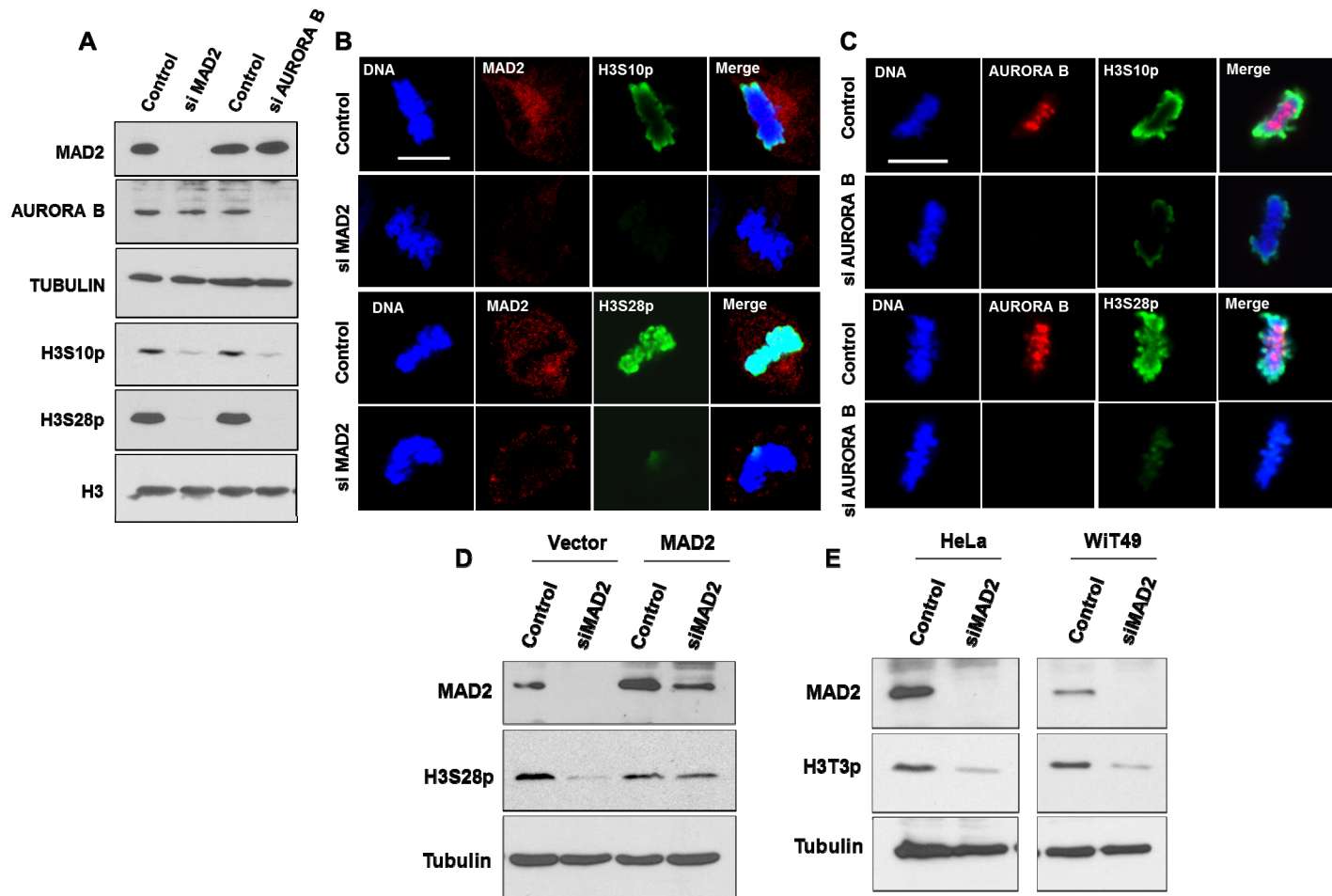
277 probed using its phospho-site specific antibody. Western blotting with histone H3 and  
278 TUBULIN antibodies were performed as controls.

279

280 **Figure 4:** MAD1 has a limited role in the regulation of AURORA B-mediated mitotic functions

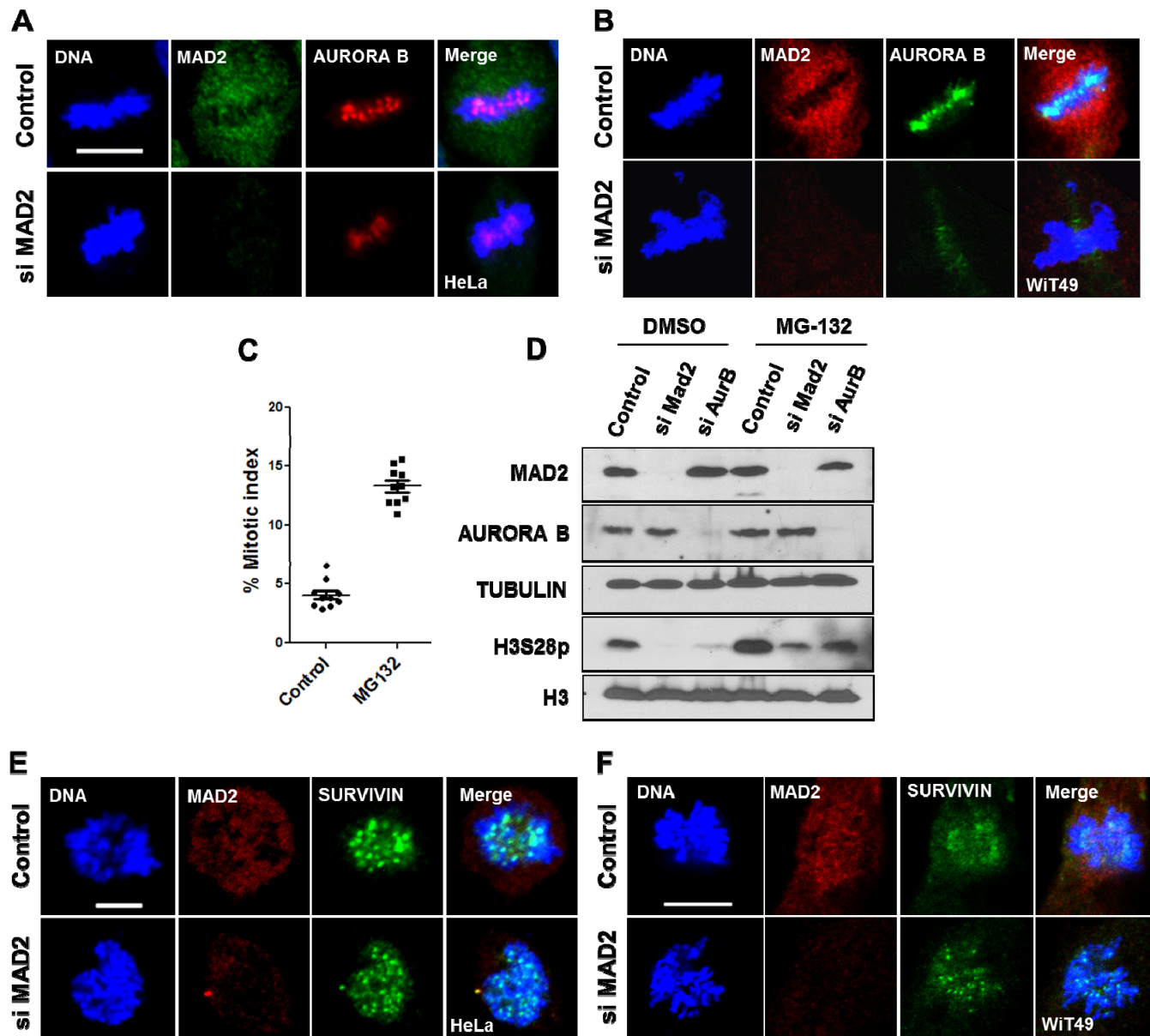
281 (A). HeLa cells were transfected with control or MAD1 siRNA for 48 hours, followed by  
282 immunoblotting with anti-MAD1 and phosphorylation specific antibodies against histone H3  
283 residues serine 10 and serine 28. Western blotting with histone H3 and TUBULIN antibodies  
284 were performed as controls. (B). Immunofluorescence analysis of control and MAD1 siRNA  
285 transfected HeLa cells was performed with the indicated antibodies to visualize Aurora B  
286 localization and the phosphorylation status of histone H3 S10 and S28 residues during  
287 metaphase. DNA was stained with Hoechst. Scale bar is 10 microns.

**Figure 1**

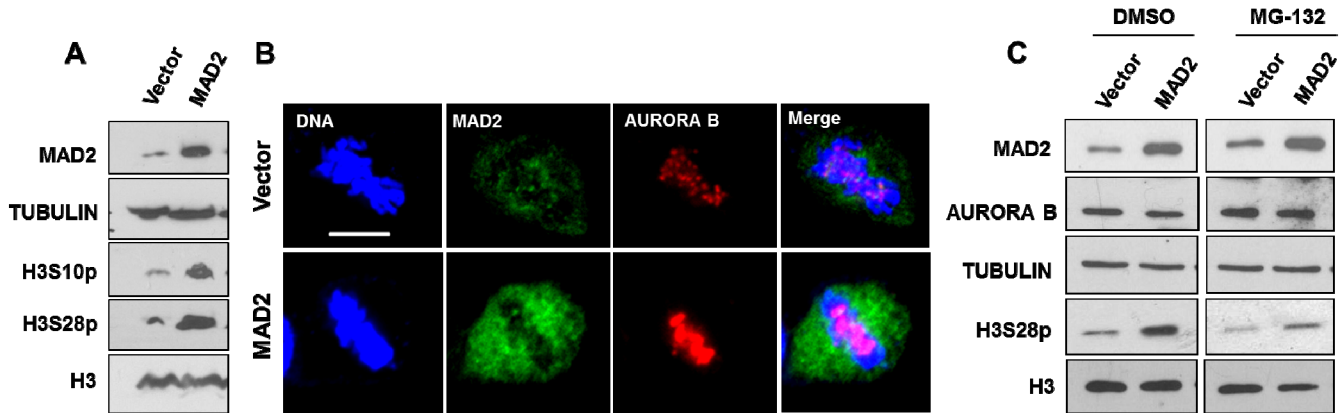




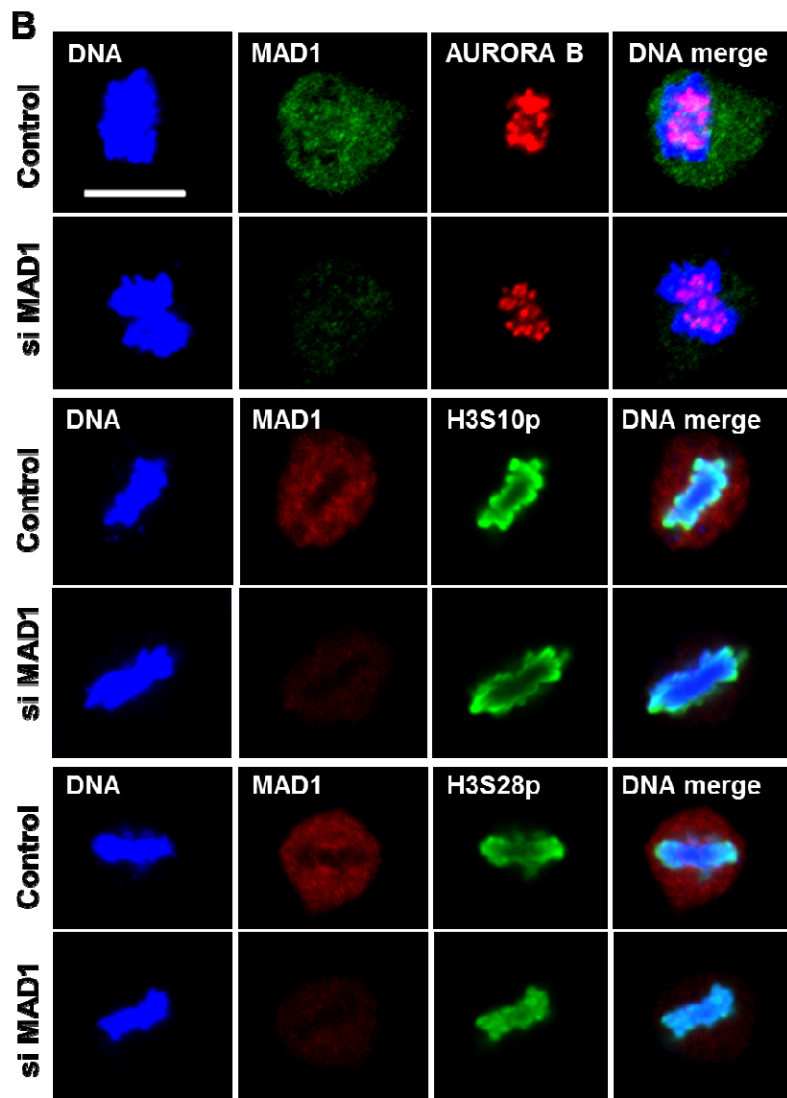
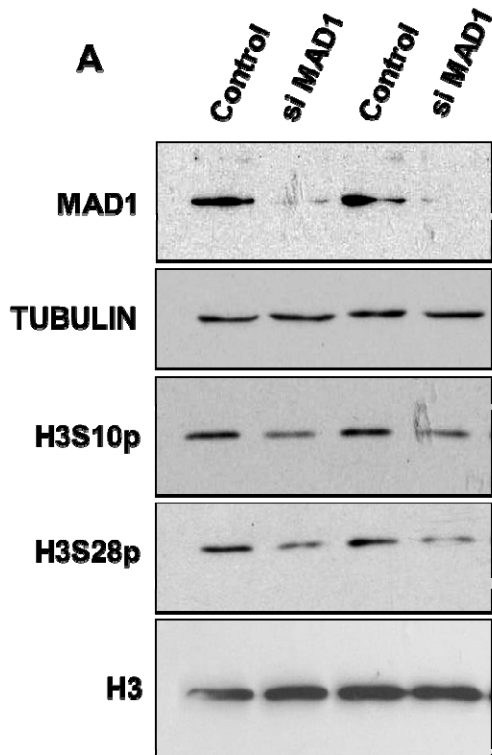
**Figure 2**



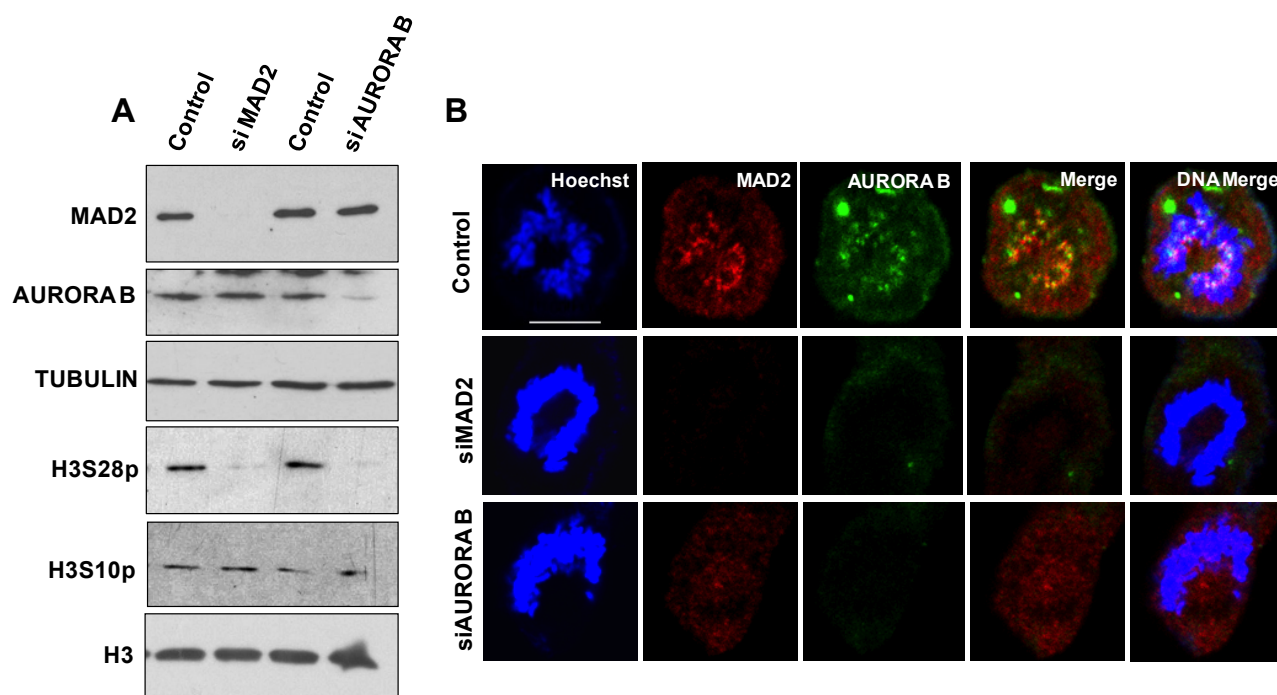
**Figure 3**



**Figure 4**



## Supplementary Figure



**Supplementary Figure S1:** MAD2 regulates Aurora B localization and the phosphorylation of histone H3 during mitosis

(A). MCF7 cells were transfected with control, MAD2 siRNA or AURORA B siRNA for 48 h, followed by immunoblotting with anti-MAD2, anti-AURORA B and anti-TUBULIN antibodies for control. The phosphorylation status of histone H3 at S10 and S28 residues was probed with its phospho-site specific antibody. Histone H3 Western blotting was performed as a control. (B). Immunofluorescence analysis of control, MAD2 and AURORA B siRNA transfected MCF7 cells was performed using MAD2 and AURORA B antibodies during metaphase. DNA was stained with Hoechst. Scale bar is 10 microns.